CLAIM AMENDMENTS

1. (currently amended): A compound of the formula

$$\mathbb{R}^3$$
 \mathbb{N}
 \mathbb{R}^3
 \mathbb{N}
 \mathbb{R}^2
 \mathbb{R}^2

and the or a pharmaceutically acceptable salts and prodrug forms salt thereof; wherein Ar represents an optionally substituted aromatic or optionally substituted heteroaromatic moiety containing 5-12 ring members wherein said heteroaromatic moiety contains one or more O, S, and/or N, 2-, 3- or 4-pyridyl, indolyl, 2- or 4-pyrimidyl, pyridazinyl, benzotriazol or benzimidazolyl,

with a proviso that optionally substituted Ar is not

wherein R⁵ is H, alkyl (1-6C), alkenyl (2-6C), alkynyl (2-6C), an aromatic or heteroaromatic moiety containing 5-11 ring members;

 $X \text{ is } NR^1, -0, \text{ or } S;$

R¹ is H, alkyl (1-8C), alkenyl (2-8C), or alkynyl (2-8C);

Z represents Nor CR4;

each of R³ and R⁴ is independently H, or a non-interfering substituent H, alkyl, alkenyl, alkynyl, acyl, aryl, alkylaryl, o-aryl, O-aryl, O-aryl, O-aroyl, NR-aryl, NR-aryl, NR-aryl, NR-aryl, NR-aryl, NR-aryl, or the hetero forms of any of the foregoing, halo, OR, NR₂, SR, -SOR, -NRSOR, -NRSO₂R, -SO₂R, -OCOR, -NRCOR, -NRCONR₂, -NRCOOR, -OCONR₂,

-COOR, -SO₃R, -CONR₂, -SO₂NR₂, -CN, -CF₃, or -NO₂, wherein each R is independently H or alkyl (1-10C);

wherein any alkyl, alkenyl, alkynyl, acyl or aryl groups contained in R³ and/or R⁴ may contain one or more heteroatoms and/or optionally be further substituted;

each R² is independently-a non-interfering substituent alkyl, alkenyl, alkynyl, acyl, aryl, alkylaryl, aroyl, O-aryl, O-aroyl, NR-aryl, NR-aryl, NR-aroyl, or the hetero forms of any of the foregoing, halo, OR, NR₂, SR, -SOR, -NRSOR, -NRSO₂R, -SO₂R, -OCOR, -NRCOR, -NRCONR₂, -NRCOOR, -OCONR₂, -COOR, -SO₃R, -CONR₂, -SO₂NR₂, -CN, -CF₃, or -NO₂, wherein each R is independently H or lower alkyl (1-4C), wherein any alkyl, alkenyl, alkynyl, acyl or aryl groups contained in R² may contain one or more heteroatoms and/or may optionally be further substituted; and

n is 0-5.

- 2. (canceled)
- 3. (canceled)
- 4. (original): The compound of claim 1, wherein the substituents on the aromatic moiety of Ar are selected from the group consisting of alkyl, alkenyl, alkynyl, acyl, aryl, alkylaryl, aroyl, O-aryl, O-aroyl, NR-aryl, NR-alkylaryl, NR-aroyl, or the hetero forms of any of the foregoing, halo, OR, NR₂, SR, -SOR, -NRSOR, -NRSO₂R, -SO₂R, -OCOR, -NRCOR, -NRCONR₂, -NRCOOR, -OCONR₂, -COOR, -SO₃R, -CONR₂, -SO₂NR₂, -CN, -CF₃, and -NO₂, wherein each R is independently H or alkyl (1-10C), and wherein any alkyl, alkenyl, alkynyl, acyl or aryl moieties contained in the substituent may contain one or more heteroatoms and/or may further be substituted by the foregoing substituents.
- 5. (currently amended): The compound of claim 1, wherein Ar is optionally substituted phenyl, 2, 3 or 4-pyridyl, indolyl, 2 or 4-pyrimidyl, or benzimidazolyl, pyridazinyl, benzotriazol or 2-pyridyl.

6. (original): The compound of claim 1, wherein n is 0-3.

7. (original): The compound of claim 1, wherein R¹ is H or lower alkyl (1-4C).

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- 8. (currently amended): The compound of claim 2 claim 1, wherein each R³ and R⁴ is independently H, alkyl (1-10C), OR, SR or NR₂ wherein R is H or alkyl (1-10C), each optionally substituted.
- 9. (original): The compound of claim 8, wherein said optional substituent is an aromatic moiety or a heterocyclic moiety, each optionally substituted.
 - 10. (original): The compound of claim 9, wherein at least one of R³ and R⁴ is H.
- 11. (currently amended): The compound of claim 3 claim 1, wherein each R² is independently alkyl, alkoxy, or halo.
 - 12. (original): The compound of claim 11, wherein each R² is independently halo.
- 13. (original): The compound of claim 4, wherein the substituents on the aromatic moiety of Ar are selected from the group consisting of alkyl, O-aryl, O-alkylaryl, NR-aryl, and N-alkylaryl wherein any alkyl or aryl contained in said substituent may further optionally be substituted.
- 14. (currently amended): The compound of claim 13, wherein said aryl aromatic moiety of Ar includes 0, 1 or 2 substituents.
- 15. (currently amended): The compound of claim 14, wherein said aryl aromatic moiety of Ar includes 0 or 1 substituents.

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16. (currently amended): The compound of elaim 2 claim 1, wherein each R³ and R⁴ is independently H, CN, COOR, OR, SR, NR₂, alkyl (1-6C), acyl (1-6C), aryl, aryloxy, arylalkyloxy, wherein R is H or alkyl (1-10C) and wherein any alkyl or aryl portions of said substituents may further be substituted with the foregoing.

- 17. (currently amended): The compound of claim 1, wherein R^{1} is H X is NH.
- 18. (currently amended): The compound of elaim 5 claim 1, wherein Ar is optionally substituted phenyl, 4-pyridyl, 3-pyridyl, 4-pyrimidyl, or 2-pyrimidyl.
- 19. (currently amended): The compound of elaim 18 claim 1, wherein Ar is optionally substituted 4-pyridyl.
- 20. (original): A method to treat conditions associated with unwanted activity of TGFβ which method comprises administering to a subject in need of such treatment an effective amount of the compound of claim 1 or a pharmaceutical composition thereof.
- 21. (currently amended): A pharmaceutical composition which comprises the compound of formula (1) claim 1 in admixture with at least one pharmaceutically acceptable excipient.
 - 22. (new) The compound of claim 17, wherein n is 1 or 2.
- 23. (new) The method of claim 20, wherein the condition associated with unwanted activity of TGF- β is a fibroproliferative disease, an autoimmune disorder, or a condition associated with eye surgery.